

REPUBLIC OF SOUTH AFRICA



REPUBLIEK VAN SUID-AFRIKA

LETTERS PATENT
(PATENTS ACT, 1978)

PATENTBRIEF
(WET OP PATENTE, 1978)

No. 96/4584

WHEREAS
NADEMAAL
UG PLUS INTERNATIONAL INC.
(Hereinafter called "the Patentee")
(Hieronder "die Patenthouer" genoem)

has applied to me for the grant of a patent in respect of an invention described and claimed in the complete specification
aansoek by my gedoen het om die verlening van 'n patent ten opsigte van 'n uitvinding wat beskryf is en waarop aanspraak
deposited at the Patent Office under the above-mentioned number, a copy of which is annexed, together with the relevant
gemaak word in die volledige spesifikasie wat by die Patentkantoor onder bovenmelde nommer ingedien is en waarvan 'n

Form P.2;
afskrif aangeheg is tesame met die betrokke Vorm P. 2;

NOW THEREFORE these Letters Patent are to grant to the Patentee a patent, the effect of which shall be to grant to the
SO IS DIT dat hierdie Patentbrief aan die Patenthouer 'n patent verleen wat die uitwerking het dat, behoudens die
Patentee in the Republic, subject to the provisions of the Act, for the duration of the patent, the right to exclude other persons
gevolgs van die Wet, aar die Patenthouer vir die duur van die patent in die Republiek die reg verleen word om ander
m making, using, exercising or disposing of the invention, so that he shall have and enjoy the whole profit and advantage
sone uit te sluit van die vervaardiging, aanwending, uitoefening of van die handsetting van die uitvinding, sodat hy al die
ruing by reason of the invention.
en voordeel wat uit die uitvinding voortspruit, verkry en geniet.

IN TESTIMONY WHEREOF the seal of the Patent Office has been affixed at Pretoria with effect from the
ER BETUIGING WAARVAN die seël van die Patentkantoor hierop te Pretoria aangebring is met ingang van die

26TH day of FEBRUARY nineteen hundred and NINETY SEVEN
dag van eenduisend negehonderd

R. J. Muller
Registrar of Patents - Registrateur van Patente

[See overleaf
Blaai om

G.P.S. 019-0127

BEST AVAILABLE COPY

REPUBLIC OF SOUTH AFRICA

LETTERS PATENT

Title

PROCESS FOR OBTAINING CHROMIUM ENRICHED
CHROMITE FROM CHROMITE ORES

APPLICANT : UG PLUS INTERNATIONAL
INC.

INVENTOR : Jean-Marc LALANCETTE,
Mario BERGERON,
François BOSSÉ and
Cyril CLERK

SHORT TITLE : Process for Obtaining
Chromium Enriched
Chromite from Chromite
Ores

PRIORITY CLAIMED : None

Law Offices of
JOHN GALGUT
23A Seventh Avenue
Parktown North
Johannesburg 2193
(011) 442-4414

JG Ref: 64974

REGISTER OF PATENTS

OFFICIAL APPLICATION NO.			LODGING DATE: PROVISIONAL		ACCEPTANCE DATE	
21	01	96 4584	22		47	1996 -12- 12
INTERNATIONAL CLASSIFICATION			LODGING DATE: COMPLETE		GRANTED DATE	
51	C 22 B		23	1996 -06- 04	1997 -02- 26	
FULL NAME(S) OF APPLICANT(S)/PATENTEE(S)						

71

UG PLUS INTERNATIONAL INC.

BEST AVAILABLE COPY

APPLICANTS SUBSTITUTED:					DATE REGISTERED	
71						

ASSIGNEE(S)					DATE REGISTERED	
71						

FULL NAME(S) OF INVENTOR(S)						
72						

Jean-Marc LALANCETTE, Mario BERGERON, François BOSSE and
Cyril CLERK

PRIORITY CLAIMED	COUNTRY		NUMBER		DATE	
N.B. Use International abbreviation for country (See Schedule 4)	33		31		32	

TITLE OF INVENTION						
54	PROCESS FOR OBTAINING CHROMIUM ENRICHED CHROMITE FROM CHROMITE ORES					

ADDRESS OF APPLICANT(S)/PATENTEE(S)						
-------------------------------------	--	--	--	--	--	--

550, Chambly, Sherbrooke, Quebec J1J 2Y2, CANADA

ADDRESS FOR SERVICE					REF:	
74	JOHN GALGUT & CO., JOHANNESBURG					64974

PATENT OF ADDITION NO.		DATE OF ANY CHANGE				
61						

FRESH APPLICATION BASED ON		DATE OF ANY CHANGE				

Patents Form P 7

REPUBLIC OF SOUTH AFRICA
PATENTS ACT, 1978
COMPLETE SPECIFICATION
[Section 30(1) – Regulation 28]

Official application No		
21	01	96145844

Lodging Date	
22	1996 -06- 04

International Classification	
51	C 22 B

Full names of Applicant(s)	
71	UG PLUS INTERNATIONAL INC.

Full name(s) of inventor(s)	
72	Jean-Marc LALANCETTE, Mario BERGERON, François BOSSÉ and Cyril CLERK

Title of invention	
54	PROCESS FOR OBTAINING CHROMIUM ENRICHED CHROMITE FROM CHROMITE ORES

TITLE OF THE INVENTION

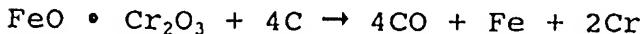
Process for obtaining chromium enriched chromite from chromite ores.

FIELD OF THE INVENTION

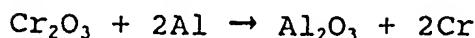
5 The present invention relates to a process for obtaining chromium enriched chromite from chromite ores or chromite concentrates.

BACKGROUND OF THE INVENTION

10 The production of metallic chromium is an important metallurgical process, this metal being an essential component in high performance alloys and in apparatuses exposed to corrosion. The elaboration of elemental chromium is made from the chromium III oxide, Cr₂O₃. If a crude form of chromium is desired, such as 15 in the ferrochrome master alloys, the chromite, FeO • Cr₂O₃, which is the most common occurrence of chromium in nature, is directly reduced with carbon in an arc furnace to give an iron-chromium alloy described by the following equation:



On the other hand, in applications where chromium must be of high purity, such as in electrochemical technologies, pure Cr₂O₃ is required as a starting material and the production of the free metal 25 is done by reaction with aluminum as described by the following equation.



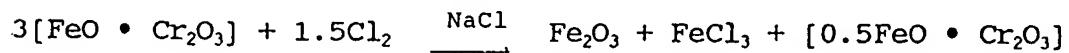
This aluminothermic reaction is quite exothermic (130 Kcal/mole) and gives fused alumina as a secondary product. Therefore, the elaboration of chromium III oxide, either as chromite of high chromium content or as purified Cr₂O₃, is an important step in the production of elemental chromium of good quality.

At present, the production of chromium III oxide consists of a rather elaborate process wherein chromite is used as the starting material. The main steps of this process consist in first fusing the chromite in the hearth of a reverberatory furnace with the addition of alkali hydroxide or carbonate and burned lime. In this way, the chromium is transformed into chromate, through the oxidizing action of the furnace flame, in the presence of an excess of oxygen. The chromate leached out with water is converted to dichromate by the addition of acid. After separation of more sparingly soluble compounds, the dichromate is deposited from the concentrated solution in a sufficiently pure state. Pure chromium III oxide is then obtained by reduction with carbon (see H. REMY, Treatise on Inorganic Chemistry, Vol. II, p. 155, Elsevier Publishing Company, 1956).

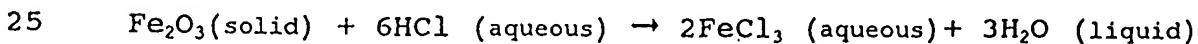
It will be readily seen that such a complex procedure makes the resulting chromium III oxide an expensive commodity.

SUMMARY OF THE INVENTION

In the course of a study of the recovery of trace amounts of platinum and related metals from chromite (see applicant's copending international application PCT/CA96/00060 filed January 30, 1996), the inventors have been led to use chlorine as a reagent for the extraction of these precious metals from the chromite. The presence of a trace amount of sodium chloride in the chromite seems to facilitate the reactions. Although chlorine turned out to be a very selective reagent for the recovery of platinoids, leaving most of the chromite unaltered, the inventors have noted that, by appropriate operating conditions, a significant amount of the iron oxide present in the chromite ($\text{FeO} \bullet \text{Cr}_2\text{O}_3$) in the presence of NaCl could be oxidized to hematite, thus increasing the Cr/Fe ratio in the residual insoluble chromite. This specific action of chlorine on chromite can be described by the following equation:



Hematite (Fe_2O_3), being readily soluble in hydrochloric acid can be removed by acid washing from the system, along with the acid soluble ferric chloride formed during the chlorination as shown by the following equation:



This relatively simple procedure, offers an interesting perspective to an easier access either to a chromite enriched in chromium.

The present invention therefore relates to a process for obtaining chromium enriched chromite from chromite ores or chromite concentrates which comprises the steps of:

- dry chlorinating the ores ore chromite concentrates at a temperature between 400°C and 750°C to produce chlorinated substrate containing acid soluble ferric oxide, acid soluble ferric chloride and acid insoluble chromium enriched chromite;
- digesting the obtained chlorinated substrate with hydrochloric acid to dissolve the ferric oxide and the ferric chloride; and
- filtering to recover said chromium enriched chromite.

In another form of the invention, the dry chlorinating step is conducted in the presence of sodium chloride.

In one form of the invention, the chromite ores are concentrated by gravimetric classification prior to chlorination.

In another form of the invention, the digesting step is carried out in the presence of heat.

Other objects and further scope of applicability of the present invention will become apparent from the detailed description given hereinafter. It should be understood, however, that this detailed description, while indicating preferred embodiments of the invention, is given by way of illustration only, since various changes and modifications within the spirit and scope of the invention will become apparent to those skilled in the art.

IN THE DRAWINGS

Figure 1 is a block diagram illustrating the various steps of one embodiment of the process of the present invention.

DESCRIPTION OF PREFERRED EMBODIMENTS

One typical chromite ore used with the present invention is that from the Bushveld complex area in South Africa, containing approximatively 10% chromium of 45% chromite wherein the ratio of Cr to Fe is about 2 to 1.

Referring to the diagram, the ore is reduced to an appropriate size by grinding. It will be readily understood that too coarse a material will prevent access to the material inside the particles and will slow down

the reaction. On the other hand, very fine grinding involves significant cost. A value of particles of about 75 microns appears as a practical compromise between reactivity and cost of grinding.

5 Alternatively, the chromite ore may be upgraded prior to chlorination so as to give an enriched concentrate containing from 30% to 65% of chromium by weight.

This ground material, either ore or 10 concentrate, is then dried to less than 0.1% of free water and heated up to the appropriate temperature for a dry chlorine treatment, sodium chloride being preferably added at this stage. This temperature of dry chlorination has been found to be variable depending on 15 the refractoriness of the ore under treatment. With certain ores, a temperature of 450°C has been sufficient. In other instances, higher temperatures (up to 700°C) have been required to achieve similar recovery. The temperature of dry chlorination is an important factor in 20 the capacity of treatment of a given apparatus. This temperature must be high enough to ensure the desired reactions while too high a temperature may prove detrimental to the equipment and may induce undesirable reactions with chromite. It has been found that a 25 temperature in the vicinity of 550°C is a preferred value. The ground ore can be heated to the selected

temperature prior to contacting with chlorine or heated up in the chlorinator itself.

The hot ore is then introduced in the dry chlorinator, which is kept at the selected temperature, and a slow stream of chlorine or chlorine containing gas is circulated through the mass at constant temperature. It must be noted that the amount of chlorine required is very small and amounts from 1.0 to 1.5 times the combined chlorine in the course of the dry chlorination. The amount of chlorine is relatively negligible, its consumption being related to the amount of reacting chromite and base metals, such as iron, nickel and copper present in the starting ore. As an indicative estimate, it can be said that some of the base metal sulfides (Cu, Ni), up to 50 percent of the iron in the chromite and less than one-tenth of one percent of the chromium in the chromite, will be converted into the corresponding soluble chlorides. In order to ensure an homogeneous distribution of the chlorine through the reacting mass, the chlorine stream can be diluted by a carrier gas, such as nitrogen for example. The dry chlorination can be achieved on a continuous basis or batch-wise by simple percolation through the mass of the reacting gas or by the fluidization of the reacting bed or by renewal of the surfaces in a rotating kiln. The contacting time required is relatively long (of the order of half an hour to two hours) depending on the nature of the ore, the

degree of completion desired, the temperature used and, to some extent, on the granulometry of the ore. For sake of simplicity of an apparatus which operates under rather aggressive conditions (chlorine at 600°C for example), a static bed of appropriate thickness has been found convenient. Since the material to be treated is to be loaded hot, the heating of the dry chlorination reactor has to be just enough to compensate for heat losses of the system.

The volatiles leaving the dry chlorinator are directed to a condensing tower and quenched by a spray of hydrochloric acid at a concentration of 6M. The temperature in this tower is of the order of 50°C and the condensable are collected and carried down by the acid spray.

The solid left in the dry chlorinator after treatment is dumped, after cooling, in a digester containing the acid from the spray in the condensing tower, and is further digested, at 100°C, for one to several hours, the stirring of the mass being ensured by recirculation of the off gases of the condensation tower through the slurry. The off gases from the digester, mostly nitrogen, hydrochloric acid vapors, chlorine and traces of sulfur compounds, are directed back to a gas treatment system after recycling condensed hydrochloric acid to the digester.

The temperature and duration of digestion in hydrochloric acid in the presence of chlorine must be adjusted in such a fashion as to ensure the complete dissolution of the soluble metal compounds. The heat increases the rate of such reactions, which could proceed nevertheless at room temperature but at a slower rate. For sake of completeness of dissolution as readily as possible of the metal, hot treatment is preferred over cold treatment and a one to several hours of contacting is retained.

Microscopic examination and diffraction analysis confirm the transformation of significant amounts of the ferrous oxide in chromite into hematite. By acid leach in hydrochloric acid, the hematite can be removed while the chromium III oxide is left undissolved. This breakdown of the spinel structure has a weakening effect on the whole crystal which is rendered more friable. The material can be used as such, being an enriched chromium ore because of FeO removal, or it can be further upgraded by taking advantage of the relatively high density of Cr₂O₃ as compared to the residual silicates initially present in the chromite ore. By application of gravimetric separation, a richer concentrate in Cr₂O₃ can be obtained.

In summary, selective oxydation of chromite can lead to the production of hematite (Fe₂O₃), ferric

chloride (FeCl_3) and chromium III oxide (Cr_2O_3) by rupture
of the spinel structure of the starting chromite. Acid
leaching by hydrochloric acid can remove selectively the
ferric oxide and chloride fraction, giving a significant
5 enrichment in Cr_2O_3 of the insoluble residual chromite
which can be further improved by gravimetric
classification or appropriate physical separation of the
gangue initially found around the chromite crystal.

10 The solubilization of the iron oxide associated
with the spinel structure of the chromite ($\text{FeO} \cdot \text{Cr}_2\text{O}_3$)
thus can be achieved to a variable extent, allowing the
removal of from five to sixty five percent of the FeO in
the starting chromite, giving an enrichment in chromite
from a ratio of 2/1 to more than 4/1.

15 The range of enrichment is a function of
temperature, duration of treatment and structure of the
starting ore.

20 The solution separated from the insoluble
chromite ore by filtration is directed to a distillation
line or concentration apparatus, such as a reverse
osmosis system, in order to recycle the hydrochloric acid
at a concentration of 6M which corresponds to the
azeotropic composition of the $\text{HCl}-\text{H}_2\text{O}$ system. The
bottoms of this distillation contain the iron, the
25 chromium and the base metals (Ni, Cu, etc...) that have

analyses of the grains which showed major signs of alterations revealed by the appearance of hematite and Cr₂O₃ crystals; 3) optical microscopy examination of the altered grains where the presence of hematite grains around the rims of the altered chromite grains could be clearly seen.

The inventors also noted that the plagioclase feldspar grains were untouched by the chlorination while the pyroxene grains showed minimal alteration, these slight alterations being related to the presence, in the starting materials, of phyllosilicates inclusions in the pyroxenes.

EXAMPLE 2

The chlorinated product obtained from Example 1 was washed with 6M hydrochloric acid. In this solution, a series of 35 elements were analysed by ICP-AES. Only iron showed significative removal from the ore by the chlorination process.

From these analysis, it can be seen that after dry chlorination and leaching with hydrochloric acid, the Cr₂O₃ is essentially intact and not dissolved. As to the iron, about 50% of this element is extracted either as the ferric chloride formed during the dry chlorination or by dissolution of the hematite. The Cr/Fe ratio is then increased to about 4/1 in the end product.

EXAMPLE 3

The chromite ore as described in Example 1 was classified in a spiral so as to give concentrates containing from 30 to 65% Cr₂O₃ by weight. These 5 concentrates, after grinding to 75 microns, were treated as in Example I with similar increases in the Cr/Fe ratio.

Although the invention has been described above with respect with one specific form, it will be evident 10 to a person skilled in the art that it may be modified and refined in various ways. For example, the grinding step may be carried out after the pre-heating step illustrated in the diagram. It is therefore wished to have it understood that the present invention should not 15 be limited in scope, except by the terms of the following claims.

The embodiments of the invention in which an exclusive property or privilege is claimed are defined as follows:

1. A process for obtaining chromium enriched chromite from chromite ores comprising the steps of:

- dry chlorinating said ores at a temperature from 400°C to 750°C to produce a chlorinated substrate containing acid soluble ferric oxide, acid soluble ferric chloride and acid insoluble chromium enriched chromite;

- digesting said chlorinated substrate with hydrochloric acid to dissolve said ferric oxide and said ferric chloride; and

- filtering to recover said chromium enriched chromite.

2. A process as defined in claim 1, further comprising the step of gravimetrically classifying said ores prior to said dry chlorinating step to obtain a chromite concentrate.

3. A process as defined in claim 1 or 2, wherein said digesting step is carried out in the presence of heat.

4. A process as defined in claim 3, wherein said digesting step is conducted in the presence of heat in the order of 100°C.

5. A process as defined in claim 4, wherein gases resulting from said digesting step are directed to a gas treatment system after recycling condensed hydrochloric acid to said digesting step.

6. A process as defined in claim 1 or 2, wherein said dry chlorinating step is conducted in the presence of sodium chloride.

7. A process as defined in claim 6, comprising the step of heating said ores prior to said dry chlorinating step.

8. A process as defined in claim 6 or 7, further comprising the step of drying said ores prior to said dry chlorinating step; said sodium chloride being introduced at said drying step.

9. A process as defined in claim 8, further comprising the step of grinding said chromite ores prior to said drying step.

10. A process as defined in claim 2, further comprising the step of grinding said chromite ores after said classifying step.

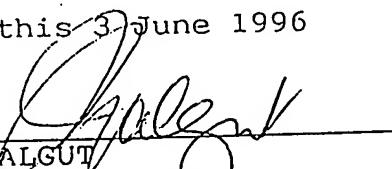
11. A process as defined in claim 9 or 10, wherein said chromite ores are ground to a size of about 75 microns.

12. A process as defined in claim 7, comprising the step of cooling said chlorinated substrate prior to said digesting step.

13. A process as defined in claim 1 or 2, wherein said temperature is about 550°C.

14. A process as defined in claim 1 or 2, further comprising the step of distillating a depleted solution resulting from said filtering step to thereby recycle the hydrochloric acid; bottoms of said distillating step being directed to a system for treating vapours from said digesting step.

Dated this 30 June 1996



JOHN GALGUT
Applicant's Patent Attorney

JG Ref: 64974